INTERNATIONAL SEARCH REPORT

International application No.

PCTAJS04/39549

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rPC(7) : GOIN 33/53 US CL : 435/7.1						
According to International Patent Classification (TPC) or to both national classification and IPC						
B. FIELDS SEARCHED						
Minimum documentation searched (classification system followed by classification symbols) U.S. : 435/7. 1						
Documentation searched other than minimum documentation Io the extent-that such documents are included in the fields searched						
Electronic data base consulted during theiinternational search (name of data bane und, where practicable:, ecurch terms used) LSPGfUB, USPAT, EPO, DERWBNT, MEDLINE, CAPLUS, SCISEARCH						
C DOCU	JMENTS CONSIDERED TO THE RELEVANT					
Category *	Citation of document, \vitl indication, where a	ppropriate,	of the relevant passages Relevant to claim?	NO.		
X	US 2003/0108937 A (WILLIAh SON et al) 31 Octo		ling. Paragruphs 1-4 and 6-19			
x	0009,001 1,001 6,0052,0057-00} (0,0096,0303,0304,00 US 2002/ 0015943 A I (BEINZ et al) 27 July 200 I 00 15,0039,0104,0043,0067,00 3,0072,0041,0005,	ei al) 27 July 200 1 filing date. Paragraphs 1-4,7.8.1 1.16-19				
	0072,0002-0005,0055			- (
Y		t al Resisiance ot'M repholino Phoaphorodiamidate Oligomers to 1,3,10 dation. Arilisense Nucleic Acid Drug Dev. 1996, Vol. 6, No. 4. pages ire document.				
Y	ANGELES et al. En*yme-linked Irπmuijosorbe π A-siiy for trkA Tyrosine Kinase Activity. 1-5 Analytical Biochemistry. \99t, Vol. 236, pages 49-55, see βmire document.					
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O Further	documents are listed in the continuation ofBox C.	D	See patent family annex.			
	pecial CUDEO 143 of oited documentiff,	Ψ _T ,	later docurns-Q published nftar thβ uit realioimi filing d to Qr ρποπτη	date		
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Date of the actual completion of the international Bearch Date of the international search report						
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Box I Observations where certain claims were found unsearchable (Continuation of Item 1 of first sheet)			
This international report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons			
1	Claim Nos because they relate to subject matter not required to be searched by this Authority, namely		
2	Claim Nos because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be earned out, specifically		
3	Claim Nos because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6 4(a)		
Box π Ob	oservations where unity of invention is lacking (Continuation of Item 2 of first sheet)		
	ional Searching Authority found multiple inventions in this international application, as follows ontinuation Sheet		
1	As all required additional search tees were timely paid by the applicant, this international search report covers all searchable claims As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee As only some of the required additional search fees were timely paid by the applicant, this, international search report covers only those claims for which tees were paid, specifically claims Nos		
4 Remark on F	No required additional search fees were timely paid by the applicant Consequently, this international search report is restπcted to the invention first mentioned in the claims, it is covered by claims Nos 1-19 Protest		

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BOX π . OBSERVATIONSWHERE UNITY OF INVENTION IS LACKING					
The inventions listed as Groups I- III do not relate to a single general inventive concept under PCT Rule 13.1 because, under PCT Rule					
13.2, they lack the same or corresponding special technical features for the following	ng reasons:				
1-19, drawn to a method of identifying a candidate beta catening modulating agent.					
2Q-22, drawn to a method of modulating pathway activity	•				
23-25, drawn to a method for diagnosing a disease					
23-23, drawn to a method for diagnosting a disease					
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The first claimed invention (claims 1-19) is drawn to a method of identifying a can	didate beta catenin modulating agent. The first				
invention fails to share a technical feature with the other claimed inventions: claim	s 20-22, drawn to a method of modulating beta catenin				
pathway with an agent that binds PLK, and claims 23-25, drawn to a method for di	agnosing a disease using a probe for PLK expression.				
Methods II and III do not share method steps with Invention 1, and PLK polypeptide	es or nucleic acids are not required as for Invention 1.				
Therefore, unity of invention is lacking.					
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Form PCT/ISA/210 (second sheet) (July 1998)